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Serial No. 10/071,809

PATENT
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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Inventor: Tinghao F. Wang
Serial No.: 10/071,809
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Title: METHOD FOR SELECTIVELY ETCHING SILICON
 AND/OR METAL SILICIDE
Examiner: DuyVu n Deo
Group Art Unit: 1765

Commissioner for Patents
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**APPLICANT'S BRIEF IN SUPPORT OF THE APPEAL TO THE BOARD OF
PATENT APPEALS AND INTERFERENCES**

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I. REAL PARTY IN INTEREST

The real party in interest is Cypress Semiconductor.

II. RELATED APPEALS AND INTERFERENCES

There are no other related appeals or interferences.

III. STATUS OF CLAIMS

All pending claims, claims 1, 3-12, 14-15, 21-23, 25, and 27, have been finally rejected and are appealed. Claims 2, 13, 16-20, 24 and 26 were previously cancelled.

IV. STATUS OF AMENDMENTS

Applicant has filed no amendments subsequent to the final rejection.

V. SUMMARY OF CLAIMED SUBJECT MATTER

WSi_x (metal silicide)/poly-Si (polysilicon) stack structures are used for gate electrodes. (Application: Page 2, Lines 11-13). Dry etching techniques are desirable that will etch a vertical profile through these stack structures without etching through the polysilicon or the gate dielectric that lies beneath the stack. (Application: Page 2, Lines 15-17). Fluorine-based etching gases have high metal silicide etching rates, but undesirably etch polysilicon and the underlying oxide that forms the gate dielectric. (Application: Page 2, Lines 19-21). Chlorine-based etching gases provide higher selectivity than fluorine-based

gases, thus preserving the polysilicon and the oxide layers, but are slower.

(Application: Page 2, Lines 22-24).

A prior study by *Nojiri* of chlorine-based etching gases showed that O₂ concentrations in the etching gas of less than 15% by volume have improved metal silicide etch rates, while O₂ concentrations of 25% by volume or more resulted in a halt in the etching process and the deposition of an undesirable film.¹ (Application: Page 3, Lines 1-6). The present invention makes use of the discovery that, contrary to these prior teachings, O₂ concentrations of at least 25% by volume in the chlorine-based etching gas, not only provide high metal silicide etch rates, but selectively etch metal silicide over polysilicon and oxides at a ratio of at least 30:1. (Application: Page 6, Lines 11-15).

Independent claim 1 provides a method of etching a metal silicide layer while fabricating an integrated circuit in a Cl₂/O₂ gas environment, where the gas includes greater than or equal to 25% O₂ by volume. (Application: Page 4, Lines 1-3). The Cl₂/O₂ gas environment is provided at a pressure of approximately 2 to 40 mili-Torr and the ratio of metal silicide to polysilicon etching is at least 30. (Application: Page 4, Lines 3-4; Page 6, Lines 13-14).

Independent claim 12 provides a method of etching a metal silicide layer while fabricating an integrated circuit in a gas environment including greater

¹ Nojiri, et al., *J. Vac. Sci. Technol.* B14(3), May/June 1996, p. 1791-1795.

than 25% O₂ by volume. (Application: Page 4, Lines 1-12). The gas environment is provided at a pressure of approximately 2 to 40 mili-Torr and the etching ratio of the metal silicide layer to an underlying polysilicon layer is at least 30. (Application: Page 4, Lines 1-12; Page 6, Lines 13-14).

Independent claim 21 provides a method of etching a metal silicide with a plasma, where the plasma includes a gas mixture including chlorine and greater than 25% O₂ by volume. (Application: Page 4, Lines 1-12; Page 7, Line 18). The etching is carried out at a pressure of approximately 2 to 40 mili-Torr and the ratio of metal silicide to polysilicon etching is at least 30. (Application: Page 4, Lines 4-5; Page 6, Lines 13-14).

VI. GROUNDS OF REJECTION TO BE REVIEWED ON APPEAL

The issues to be decided on this appeal are as follows:

Whether claims 1, 3-12, 14-15, 21-23, 25, and 27 are obvious under 35 U.S.C. § 103(a) over U.S. Pat. No. 5,880,033 to Tsai (*Tsai*) alone, or in view of *Tsai* in combination with U.S. Pat. No. 6,150,250 to Tabara et al. (*Tabara*), or in view of *Tsai* and *Tabara* in combination with Langley et al., *Semiconductor International*, October 1989 (*Langley*).

VII. ARGUMENT

The Examiner has failed to establish a *prima facie* of obviousness for any of the pending independent claims. In particular, each of independent claims 1, 12, and 21 include an etching selective enough to provide a metal silicide to polysilicon etch selectivity of at least 30. The closest metal silicide to polysilicon etching selectivity that may be found in the cited references is 5, found in *Tsai*. (Col. 9, Line 45). Thus, the present invention provides an etching selectivity improvement of at least six times that found in the cited art.

Furthermore, *Tsai* teaches away from the present invention, teaching that increasing the oxygen concentration of the etching gas provides an increase in the rate of polysilicon etching and a concurrent decrease in etching selectivity.²

Tabara and *Langley* fail to cure the deficiencies of *Tsai*.

² This is true even though a drafting error resulted in an obvious misdescription of Figures 4 and 5 that states that etching selectivity increased with increasing O₂. (Col. 7, Lines 64-66; Col. 8, Lines 5-7).

A. The Cited References Fail to Render Obvious Under 35 U.S.C. § 103 an Etching Process Providing a Metal Silicide to Polysilicon Etch Selectivity of at Least 30.

1. Independent claims 1, 12, and 21 include a metal silicide to polysilicon etch selectivity of at least 30, a value not provided, suggested, or obtainable from the technology described in the cited references.

To establish a *prima facie* case of obviousness, in addition to a motivation to combine and an expectation of success, the cited art must teach or suggest every element of the claims. M.P.E.P. § 2143 citing *In re Vaeck*, 947 F.2d 488, 20 USPQ2d 1438 (Fed. Cir. 1991). Even though *Tsai* is directed to the same goals as the present application, providing a high etching rate while selectivity etching a metal silicide over polysilicon, the best selectivity obtained in *Tsai* was a 5:1 etching ratio of metal silicide to polysilicon. (Col. 9, Line 45). Thus, *Tsai* cannot teach or suggest the claimed etching selectivity of at least 30 (same as a 30:1 ratio).

If the teachings of *Tsai* could have made the presently claimed selectivity value of 30 obvious, *Tsai* would have obtained this higher etching selectivity because the reference is directed to same problem as Applicant's invention, high etching selectivity. Instead of being obvious over *Tsai*'s selectivity ratio of 5, the at least 30 etching ratio of present claims 1, 12, and 21 provides a substantial improvement, a six time increase in etching selectivity.

Tsai could not have obtained etching selectivity ratios higher than 5 at least because the highest concentration of O₂ used in the etchant gas was approximately 14% of the total gas volume. (Col. 3, Lines 25-28). In the Examples, only 3% O₂ was actually used. (Col. 9, Lines 24-26). The O₂ percentages of *Tsai* are substantially lower than those of the present claims, where at least 25% O₂ is indicated. The present claims use a significantly higher concentration of O₂ in the etchant gas to obtain the markedly higher etching selectivity of at least 30.

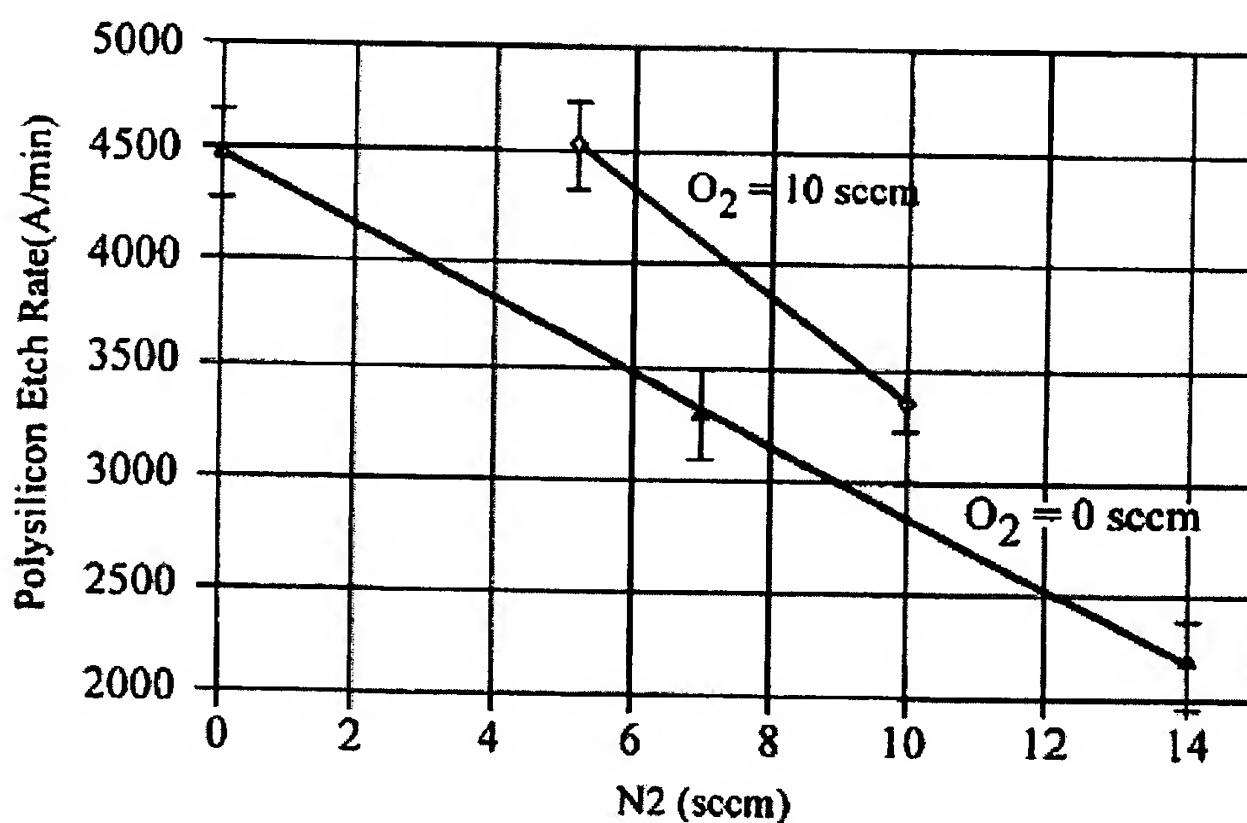
Neither could *Tsai* have suggested that Applicant's higher percentage of O₂ in the etchant gas could increase etching selectivity to provide the at least 30 value of the present claims for at least two reasons. First, *Tsai* explicitly teaches away from higher O₂ concentrations, stating that "excessively high³ flow rates of the oxygen gas can cause more isotropic etching of the substrate ... and can also result in excessively low dielectric etch rates." (Col. 7, Lines 15-20).

Furthermore, this statement from *Tsai* is consistent with the prior teachings of *Nojiri*, which were accepted when *Tsai* was filed and before the unexpected results of the present invention were obtained.

Second, *Tsai* could not suggest the at least 30 selectivity value of the present claims because the experimental data presented in *Tsai* also teaches

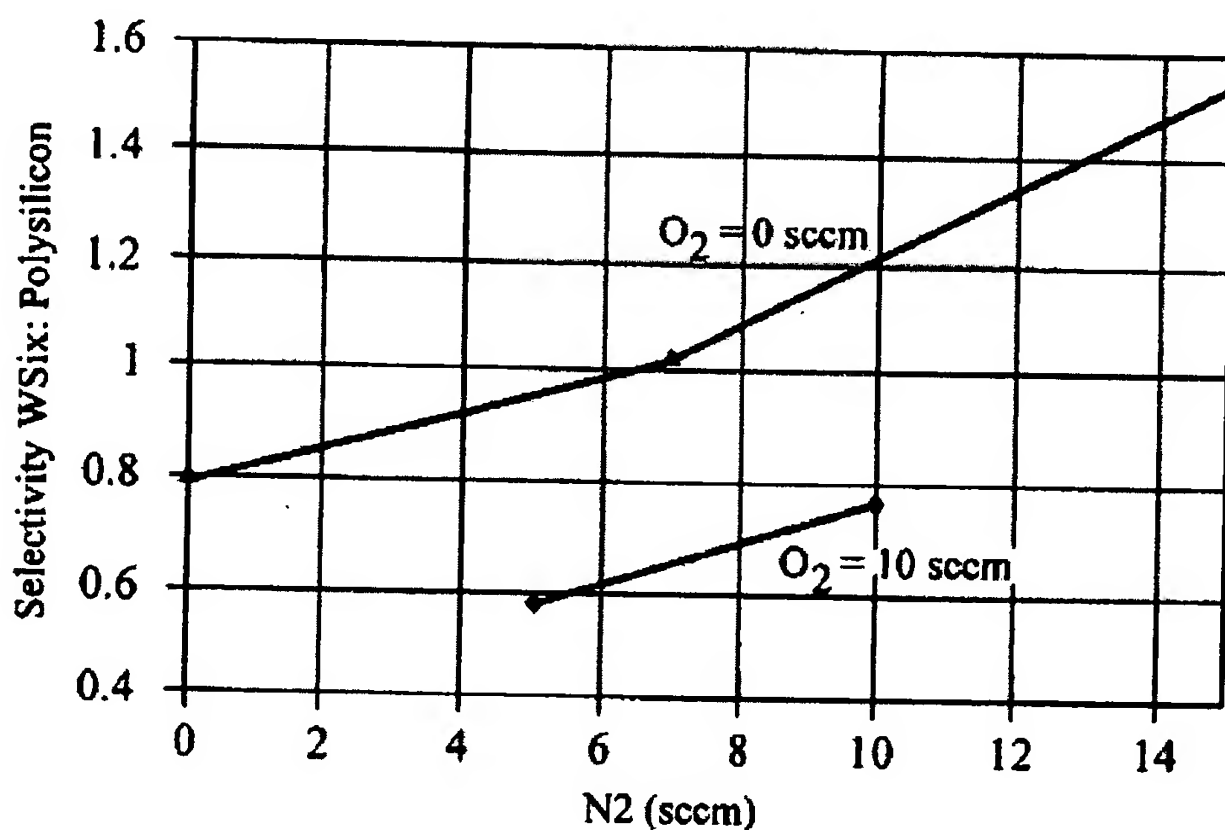
³ While "excessively high" may not be numerically specific, one of ordinary skill in the art would understand the presently claimed value of at least 25% to be "excessively higher" than the 3% to 7% values used in *Tsai*.

away from Applicant's use of high O₂ concentration to provide significantly increased selectivity. Figure 3 of *Tsai* establishes that when the O₂ concentration of the etchant gas is increased from 0 to about 7%, a beneficial increase in the rate of metal silicide etching results. However, Figure 4, reproduced below, shows that the same ~7% increase in the O₂ concentration of the etchant gas has the undesirable side effect of concurrently increasing the rate of polysilicon etching. In fact, when the O₂ concentration of the etching gas was increased from 0 to about 7%, the rate of polysilicon increased from about 3250 to about 4000 A/min.⁴



⁴ These values are at the ~7 sccm N₂ value; however, they are not significantly different at other N₂ flow rates.

Figure 5, reproduced below, compares the rates of metal silicide etching from Figures 3 to the rates of polysilicon etching from Figure 4 as selectivity ratios, with higher selectivity values representing enhanced metal silicide to polysilicon etching. As seen in Figure 5, when the O₂ concentration of the etching gas was increased from 0 to about 7%, the selectivity ratio decreased from about 1 to about 0.65 (an approximately 35% decrease).⁵



The data of *Tsai* show that an increase in the O₂ concentration of the etching gas increases the rate of metal silicide etching, while concurrently reducing the selectivity of metal silicide to polysilicon etching. Not a single data

⁵ These values are at the ~7 sccm N₂ value; however, they are not significantly different at other N₂ flow rates.

point obtained with the etching gas containing ~7% O₂ had an etching selectivity greater than that obtained with the etching gas lacking O₂.⁶

Tsai does not disclose and cannot suggest the use of etching gases having higher O₂ concentrations to obtain enhanced etching selectivity because the data only show a decrease in etching selectivity with higher O₂ content. *Tsai* cannot suggest the opposite of what it describes and can only teach away from claims 1, 12, and 21, where a 25% or greater concentration of O₂ in the etchant gas provided an etching selectivity of at least 30.

The Examiner has failed to establish a *prima facie* case of obviousness because *Tsai* fails to disclose or suggest that an etching selectivity of at least 30 (a 6 time increase over the highest obtained by *Tsai*) could be obtained by increasing the O₂ concentration of the etchant gas from the 3% of *Tsai* to greater than or equal to 25%. When considered as a whole, *Tsai* teaches that increasing the O₂ concentration of a chlorine-based etchant gas provides a desirable increase in the rate of metal silicide etching while concurrently providing an undesirable decrease in metal silicide to polysilicon etching that may be partially

⁶ The Applicant would like to point out that with regard to Figures 4 and 5, *Tsai* states that "the polysilicon etch rate is lowered for increasing flow rates of O₂ as well as increasing flow rates of N₂" (Fig. 4) and that "the etching selectivity ratios increases [sic] for both increasing flow rates of O₂ and increasing flow rates of N₂" (Fig. 5). (Col. 7, lines 64-66; Col. 8, lines 5-7). However, it is clear from the Figures that a drafting error occurred with regards to the "increasing flow rates of O₂" in both passages, which should read "**decreasing** flow rates of O₂." The graphs may not be interpreted in any other manner.

counteracted by adding N₂ gas to the etchant gas.⁷ *Tsai* does not describe or suggest the selectivity and O₂ concentrations indicated in present claims 1, 12, and 21. Thus, the rejections over *Tsai* under 35 U.S.C. § 103 must be removed.

The combination of *Tsai* with *Tabara* also fails to establish a *prima facie* case of obviousness with regard to claims 1, 12, and 21 because like *Tsai*, *Tabara* teaches that regardless of the O₂ concentration of the etchant gas, polysilicon is always etched faster than metal silicide. *Tabara* describes the etching of WSi₂ or polysilicon using TiN or TiON as an etching mask. The purpose of *Tabara* is to provide etching conditions that selectively etch the metal silicide or polysilicon layer with respect to the mask. (Col. 2, Lines 49-54). Very similar etching conditions are used to etch both WSi₂ and polysilicon with Cl₂/O₂ etching gases that include from about 31% to about 38% O₂. (Col. 7, lines 9-26).

Figure 20 of *Tabara* establishes that for any attempted O₂ flow rate the etch selectivity of Si/TiN is always greater than the selectivity of WSi₂/TiN.⁸ Thus, polysilicon is always etched faster than the WSi₂ metal silicide. *Tabara* provides no suggestion that greater metal silicide etching selectivity, certainly

⁷ A reference must be treated for all it teaches. W.L. Gore & Assocs. v. Garlock, Inc., 721 F.2d 1540, 1551, 220 U.S.P.Q. 303, 312-13 (Fed. Cir. 1983) (stating in determining obviousness, the invention must be considered as a whole without the benefit of hindsight of Applicant's disclosure).

⁸ Compare the curve on the upper left with the curve at the bottom; all other curves describe irrelevant WSi₂/TiON etch selection ratios.

not the at least 30 value of present claims 1, 12, and 21, may be obtained for a metal silicide/polysilicon etch. In fact, *Tabara* teaches away from the present invention by showing that regardless of the O₂ concentration, Cl₂/O₂ etchant gases cannot selectively etch a metal silicide over a polysilicon.

Tabara fails to provide *Tsai* with the missing etching selectivity of at least 30, which is an element necessary to establish a *prima facie* case of obviousness. Thus, the 35 U.S.C. § 103 rejections of claims 1, 12, and 21 over *Tsai* in combination with *Tabara* must be withdrawn.

The combination of *Tsai* with *Langley* or *Tabara* with *Langley* also fails to establish a *prima facie* case of obviousness. *Langley* was cited for describing a breakthrough etching using CF₄, a concept present in Applicant's dependent claims 22, 23, and 25. However, *Langley* provides no suggestion regarding how to etch a metal silicide with respect to a polysilicon with a selectivity ratio of at least 30 during a metal silicide etch. Thus, the combination of *Langley* with *Tsai* or *Tabara* cannot establish a *prima facie* case of obviousness with regard to present claims 1, 12, and 21 and the rejections under 35 U.S.C. § 103 must be withdrawn.

2. In addition to the selectivity ratio of at least 30 and the 25% O₂ by volume elements of independent claims 1, 12, and 21, independent claim 1 further indicates that the etchant gas includes Cl₂ and O₂.

A *prima facie* case of obviousness also may not be established for claim 1, because the claim is differentiated from *Tsai* through the exclusion of N₂ gas. To establish a *prima facie* case of obviousness, an expectation of success must be found in the cited reference, not Applicant's disclosure. M.P.E.P. § 2143 citing In re Vaeck.

As stated above, *Tsai* teaches that while an increased concentration of O₂ in the etchant gas increases the rate of metal silicide and polysilicon leading to inferior etching selectivity, the selectivity may be improved while maintaining the increased overall etching rate by adding N₂ gas to the mixture. *Tsai* makes this clear by stating that "[t]he nitrogen gas provides unexpected results in combination with chlorine and oxygen gases" and that "[i]ncreasing the nitrogen gas significantly lowers the rate of etching of polysilicon layer 24, without [adversely] affecting the rate of etching of the metal silicide layer 22." (Col. 7, Lines 25-26; Col. 7, Lines 26-29).

Based on the teachings of *Tsai*, one could never expect that a substantial increase in the O₂ concentration of the etchant gas in the absence of N₂ could successfully provide any increase in etching selectivity, certainly not the at least 30 etching ratio of Applicant's claim 1. *Tabara* and *Langley* fail to meaningfully

address metal silicide/polysilicon etching selectivity and, thus, cannot cure the deficiencies of *Tsai*.

Without relying on Applicant's disclosure, the Examiner failed to establish any expectation that one could successfully increase etching selectivity by increasing the O₂ concentration of the etchant gas. Thus, *Tsai*, *Tabara*, and *Langley*, alone or in combination, cannot establish a *prima facie* case of obviousness with regard to claim 1, and the rejections under 35 U.S.C. § 103 must be withdrawn.


VIII. CONCLUSION

For the foregoing reasons, the claim rejections applied by the Examiner under 35 U.S.C. § 103 are unsustainable. Applicants respectfully request reversal of the Examiner's rejections.

Respectfully submitted,

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IX. CLAIMS APPENDIX

1. (Previously Presented) A method comprising, etching a metal silicide layer during fabrication of an integrated circuit in a Cl_2/O_2 environment having an O_2 concentration of greater than or equal to 25% by volume,
wherein the Cl_2/O_2 environment is provided at a pressure of approximately 2-40 mili-Torr, and wherein the etching is a metal silicide etch that is selective to poly-silicon with a ratio of etch rates of at least 30.
2. (Cancelled)
3. (Original) The method of claim 2 wherein the pressure is approximately 3 mili-Torr.
4. (Original) The method of claim 1 wherein the Cl_2/O_2 environment is provided in a reactor with a source power of approximately 200 - 2000 Watts.
5. (Original) The method of claim 4 wherein the source power is approximately 400 Watts.
6. (Original) The method of claim 1 wherein the Cl_2/O_2 environment is provided in a reactor having a bias power of approximately 35 to 400 Watts.
7. (Original) The method of claim 6 wherein the reactor has a bias power of approximately 50 Watts.

8. (Original) The method of claim 1 wherein the metal silicide layer is a tungsten silicide layer.
9. (Original) The method of claim 1 wherein the Cl_2/O_2 environment comprises approximately 45 sccm Cl_2 and 30 sccm O_2 .
10. (Original) The method of claim 9 wherein the Cl_2/O_2 environment is provided for a time period sufficient to completely etch the metal silicide layer.
11. (Original) The method of claim 9 wherein the time period is approximately 30 seconds.
12. (Previously Presented) A method comprising etching a metal silicide layer during fabrication of an integrated circuit in an environment having a concentration of O_2 greater than 25% by volume so as to selectively etch the metal silicide layer with respect to an underlying poly-silicon layer with a ratio of etch rates of at least 30,
wherein the etching is carried out at a pressure of 2-40 mili-Torr.
13. (Cancelled)
14. (Original) The method of claim 12 wherein the environment comprises approximately 45 sccm Cl_2 and 30 sccm O_2 .
15. (Original) The method of claim 12 wherein the metal silicide is chosen from the group consisting of tungsten silicide, chromium silicide and titanium silicide.

16-20. (Cancelled)

21. (Previously Presented) A method of etching a metal silicide, comprising etching of the metal silicide with a plasma, wherein the plasma is prepared from a gas mixture comprising: chlorine, and greater than 25% by volume oxygen, the etching is carried out at a pressure of 2-40 mili-Torr, and the etching is a metal silicide etch that is selective to poly-silicon with a ratio of etch rates of at least 30.

22. (Previously Presented) The method of claim 21, further comprising, prior to said etching, a breakthrough etch.

23. (Previously Presented) The method of claim 22, wherein said breakthrough etch comprises etching with a plasma prepared from a gas comprising CF_4 .

24. (Cancelled)

25. (Previously Presented) The method of claim 1, further comprising, prior to said etching, a breakthrough etch.

26. (Cancelled)

27. (Previously Presented) The method of claim 21, wherein said gas mixture comprises: chlorine and from 25% to 30% by volume oxygen.